



(86) Date de dépôt PCT/PCT Filing Date: 2013/01/18
(87) Date publication PCT/PCT Publication Date: 2013/07/25
(85) Entrée phase nationale/National Entry: 2014/07/16
(86) N° demande PCT/PCT Application No.: US 2013/022041
(87) N° publication PCT/PCT Publication No.: 2013/109815
(30) Priorité/Priority: 2012/01/19 (US61/588,437)

(51) Cl.Int./Int.Cl. *A61B 5/0215* (2006.01)
(71) Demandeur/Applicant:
VOLCANO CORPORATION, US
(72) Inventeurs/Inventors:
DORANDO, DALE GENE, US;
WONG, DAVID, US;
ALPERT, HOWARD DAVID, US
(74) Agent: GOWLING LAFLEUR HENDERSON LLP

(54) Titre : DISPOSITIFS, SYSTEMES ET PROCEDES D'INTERFACE UTILISABLES AVEC DES DISPOSITIFS DE SURVEILLANCE DE LA PRESSION INTRAVASCULAIRE
(54) Title: INTERFACE DEVICES, SYSTEMS, AND METHODS FOR USE WITH INTRAVASCULAR PRESSURE MONITORING DEVICES

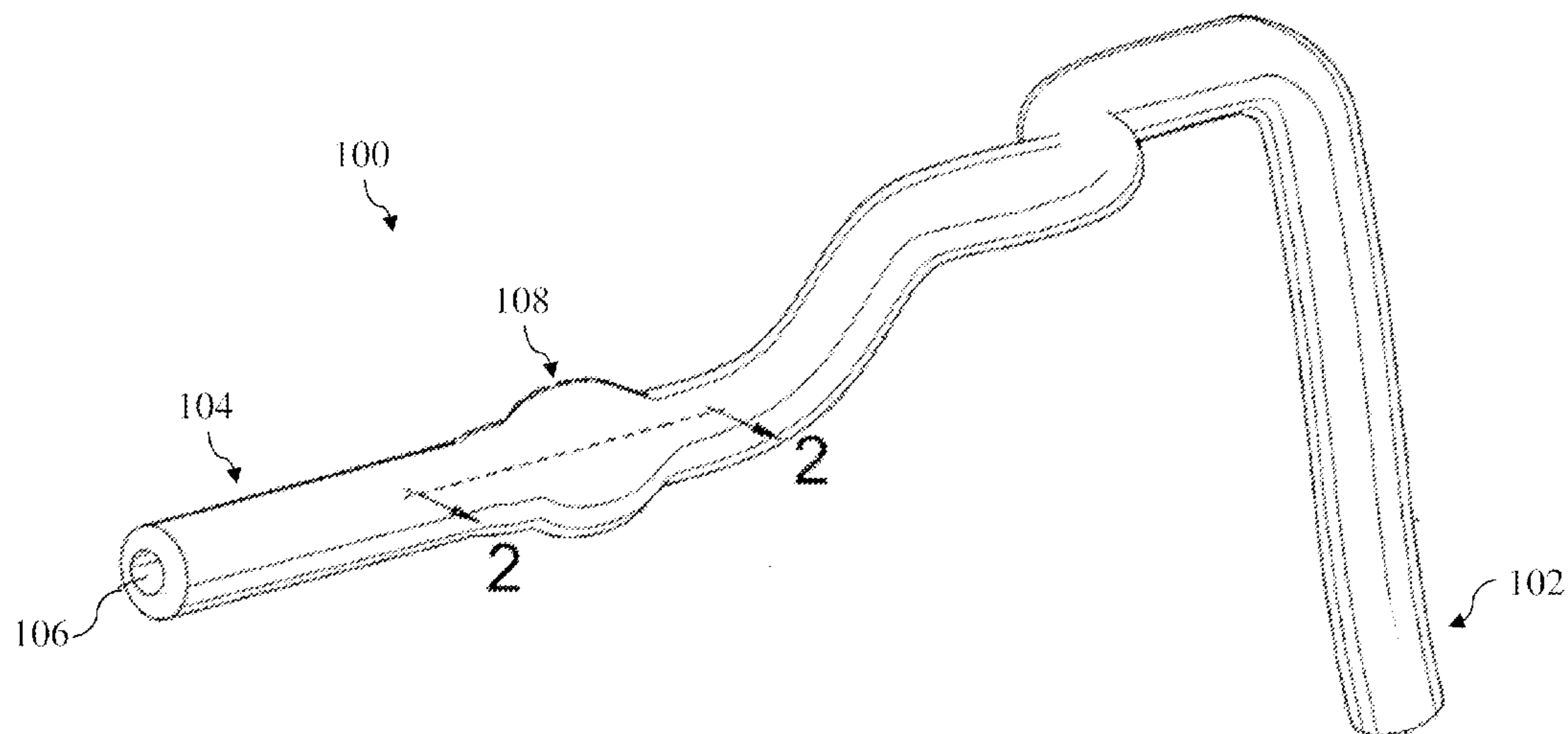


Fig. 1

(57) **Abrégé/Abstract:**

Embodiments of the present disclosure are configured to assess the severity of a blockage in a vessel and, in particular, a stenosis in a blood vessel. In some particular embodiments, the devices, systems, and methods of the present disclosure are configured to provide FFR measurements in a small, compact device that integrates with existing proximal and distal pressure measurement systems and does not require a separate power source.



(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property
Organization
International Bureau



(10) International Publication Number
WO 2013/109815 A1

(43) International Publication Date
25 July 2013 (25.07.2013)

(51) International Patent Classification:
A61B 5/0215 (2006.01)

(21) International Application Number:
PCT/US2013/022041

(22) International Filing Date:
18 January 2013 (18.01.2013)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
61/588,437 19 January 2012 (19.01.2012) US

(71) Applicant: **VOLCANO CORPORATION** [US/US];
3661 Valley Centre Drive, Suite 200, San Diego, California 92130 (US).

(72) Inventors; and

(71) Applicants (for US only): **DORANDO, Dale Gene** [US/US]; 3521 Cothrin Ranch Road, Shingle Springs, California 95682 (US). **WONG, David** [US/US]; 5539 Rainbow Circle, Santa Rosa, California 95409 (US). **ALPERT, Howard David** [US/US]; 1596 Zapata Drive, El Dorado Hills, California 95762 (US).

(74) Agents: **JOHNSON, Dustin T.** et al.; Haynes and Boone, LLP, IP Section, 2323 Victory Avenue, Suite 700, Dallas, Texas 75219 (US).

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

- with international search report (Art. 21(3))
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments (Rule 48.2(h))

(54) Title: INTERFACE DEVICES, SYSTEMS, AND METHODS FOR USE WITH INTRAVASCULAR PRESSURE MONITORING DEVICES

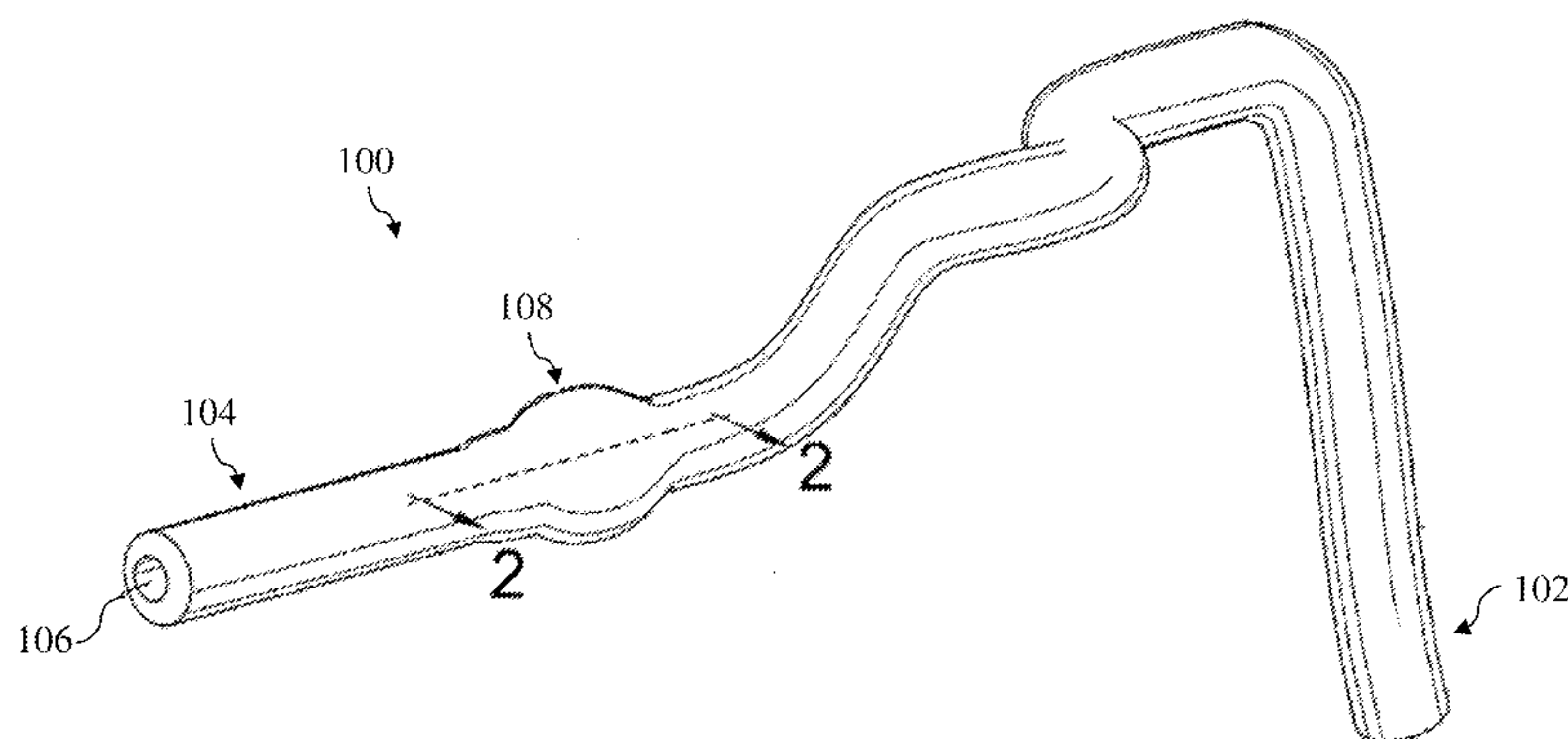


Fig. 1

(57) Abstract: Embodiments of the present disclosure are configured to assess the severity of a blockage in a vessel and, in particular, a stenosis in a blood vessel. In some particular embodiments, the devices, systems, and methods of the present disclosure are configured to provide FFR measurements in a small, compact device that integrates with existing proximal and distal pressure measurement systems and does not require a separate power source.

WO 2013/109815 A1

INTERFACE DEVICES, SYSTEMS, AND METHODS FOR USE WITH
INTRAVASCULAR PRESSURE MONITORING DEVICES

5 TECHNICAL FIELD

The present disclosure relates generally to the assessment of vessels and, in particular, the assessment of the severity of a blockage or other restriction to the flow of fluid through a vessel. Aspects of the present disclosure are particularly suited for evaluation of biological vessels in some instances. For example, some particular embodiments of the present
10 disclosure are specifically configured for the evaluation of a stenosis of a human blood vessel.

BACKGROUND

A currently accepted technique for assessing the severity of a stenosis in a blood
15 vessel, including ischemia causing lesions, is fractional flow reserve (FFR). FFR is a calculation of the ratio of a distal pressure measurement (taken on the distal side of the stenosis) relative to a proximal pressure measurement (taken on the proximal side of the stenosis). FFR provides an index of stenosis severity that allows determination as to whether the blockage limits blood flow within the vessel to an extent that treatment is required. The
20 normal value of FFR in a healthy vessel is 1.00, while values less than about 0.80 are generally deemed significant and require treatment. Common treatment options include angioplasty and stenting.

Coronary blood flow is unique in that it is affected not only by fluctuations in the pressure arising proximally (as in the aorta) but is also simultaneously affected by
25 fluctuations arising distally in the microcirculation. Accordingly, it is not possible to accurately assess the severity of a coronary stenosis by simply measuring the fall in mean or peak pressure across the stenosis because the distal coronary pressure is not purely a residual of the pressure transmitted from the aortic end of the vessel. As a result, for an effective calculation of FFR within the coronary arteries, it is necessary to reduce the vascular
30 resistance within the vessel. Currently, pharmacological hyperemic agents, such as adenosine, are administered to reduce and stabilize the resistance within the coronary arteries. These potent vasodilator agents reduce the dramatic fluctuation in resistance (predominantly by reducing the microcirculation resistance associated with the systolic portion of the heart cycle) to obtain a relatively stable and minimal resistance value.

However, the administration of hyperemic agents is not always possible or advisable. First, the clinical effort of administering hyperemic agents can be significant. In some countries (particularly the United States), hyperemic agents such as adenosine are expensive, and time consuming to obtain when delivered intravenously (IV). In that regard, IV-

5 delivered adenosine is generally mixed on a case-by-case basis in the hospital pharmacy. It can take a significant amount of time and effort to get the adenosine prepared and delivered to the operating area. These logistic hurdles can impact a physician's decision to use FFR. Second, some patients have contraindications to the use of hyperemic agents such as asthma, severe COPD, hypotension, bradycardia, low cardiac ejection fraction, recent myocardial

10 infarction, and/or other factors that prevent the administration of hyperemic agents. Third, many patients find the administration of hyperemic agents to be uncomfortable, which is only compounded by the fact that the hyperemic agent may need to be applied multiple times during the course of a procedure to obtain FFR measurements. Fourth, the administration of a hyperemic agent may also require central venous access (*e.g.*, a central venous sheath) that

15 might otherwise be avoided. Finally, not all patients respond as expected to hyperemic agents and, in some instances, it is difficult to identify these patients before administration of the hyperemic agent.

To obtain FFR measurements, one or more ultra-miniature sensors placed on the distal portion of a flexible device, such as a catheter or guide wire used for catheterization

20 procedures, are utilized to obtain the distal pressure measurement, while a sensor connected to a measurement instrument, often called the hemodynamic system, is utilized to obtain the proximal or aortic pressure measurement. Currently only large expensive systems or a combination of multiple devices connected to the distal pressure wire and the hemodynamic system can calculate and display an FFR measurement. In that regard, to calculate the FFR

25 these devices require both the aortic or proximal pressure measurement and the coronary artery or distal pressure measurement. Accordingly, these systems require the catheter lab's hemodynamic system to have a high level analog voltage output. "High level" in this context generally implies 100mmHg/Volt output. Unfortunately, there are many hemodynamic systems that don't provide a high level output. As a result, when using these hemodynamic

30 systems, providing an FFR measurement is difficult if not impossible. Further, space in a typical catheter lab is extremely limited. Consequently, devices that are large and located in the catheter lab are disfavored compared to smaller derives, especially if the smaller device can provide much if not all of the functionality of the larger device. As a result, it is highly

desirable to have a device that that can display FFR and yet is small and lightweight that can sit on, or near, the patient bed and be easily read by the physician.

Further, most pressure measurement devices require an extra source of power like an AC adapter or wall plug. This adds to wire clutter and available medical grade AC outlets are
5 not often available near the patient bed. In addition, any device that uses AC power must undergo stringent safety precautions to reduce patient risk due to leakage currents. Batteries are another alternative for power. But, batteries must be replaced, disposed of correctly and have a finite shelf life.

Accordingly, there remains a need for improved devices, systems, and methods for
10 assessing the severity of a blockage in a vessel and, in particular, a stenosis in a blood vessel. In that regard, there remains a need for improved devices, systems, and methods for providing FFR measurements that have a small, compact size (*e.g.*, suitable for hand-held use), integrate with existing proximal and distal pressure measurement devices, and do not require a separate power source.

15

SUMMARY

Embodiments of the present disclosure are configured to assess the severity of a blockage in a vessel and, in particular, a stenosis in a blood vessel. In some particular embodiments, the devices, systems, and methods of the present disclosure are configured to provide FFR measurements in a small, compact device that integrates with existing proximal and distal pressure measurement systems and does not require a separate power source.

In one embodiment, an interface for intravascular pressure sensing devices is provided. The interface comprises: a distal input configured to receive a distal pressure signal from a distal pressure sensing device; a distal output configured to output the distal pressure signal to a hemodynamic system in a format useable by the hemodynamic system; a proximal input configured to receive a proximal pressure signal from a proximal pressure sensing device; a proximal output configured to output the proximal pressure signal to the hemodynamic system in a format useable by the hemodynamic system; a processor coupled to the distal input, distal output, proximal input, and proximal output, the processor configured to calculate a pressure differential between the distal pressure and the proximal pressure based on the received distal pressure signal and the received proximal pressure signal; and a display in communication with the processor and configured to display the pressure differential calculated by the processor. In some embodiments, the distal input, distal output, proximal input, proximal output, processor, and display are secured to a housing. Further, in some instances the distal pressure sensing device is a pressure-sensing guidewire and the proximal pressure sensing device is a pressure-sensing catheter configured for use with the hemodynamic system.

In another embodiment, a system for evaluating a vascular stenosis is provided. The system comprises: a distal pressure sensing device sized and shaped for insertion into human vasculature; a proximal pressure sensing device sized and shaped for insertion into human vasculature; and an interface, where the interface includes: a distal input configured to receive a distal pressure signal from the distal pressure sensing device; a proximal input configured to receive a proximal pressure signal from the proximal pressure sensing device; a proximal output configured to output the proximal pressure signal to a processing system in a format useable by the processing system; a processor coupled to the distal input, proximal input, and proximal output, the processor configured to calculate a pressure differential between the distal pressure and the proximal pressure based on the received distal pressure signal and the received proximal pressure signal; and a display in communication with the processor and configured to display the pressure differential calculated by the processor. In

some instances, the processing system is a hemodynamic system. Further, in some instances the distal input, distal output, proximal input, proximal output, processor, and display are secured to a housing. In some instances, the distal pressure sensing device is a pressure-sensing guidewire and the proximal pressure sensing device is a pressure-sensing catheter
5 configured for use with the processing system.

Additional aspects, features, and advantages of the present disclosure will become apparent from the following detailed description.

BRIEF DESCRIPTION OF THE DRAWINGS

Illustrative embodiments of the present disclosure will be described with reference to the accompanying drawings, of which:

FIG. 1 is a diagrammatic perspective view of a vessel having a stenosis according to
5 an embodiment of the present disclosure.

FIG. 2 is a diagrammatic, partial cross-sectional perspective view of a portion of the vessel of Fig. 1 taken along section line 2-2 of Fig. 1.

FIG. 3 is a diagrammatic, partial cross-sectional perspective view of the vessel of Figs. 1 and 2 with instruments positioned therein according to an embodiment of the present
10 disclosure.

FIG. 4 is a diagrammatic, schematic view of a system according to an embodiment of the present disclosure.

FIG. 5 is a diagrammatic, schematic view of an interface device of the system of Fig. 4 according to an embodiment of the present disclosure.

FIG. 6 is a diagrammatic, schematic view of a portion of the interface device of Fig. 5
15 according to an embodiment of the present disclosure.

FIG. 7 is a diagrammatic, schematic view of a portion of the interface device of Fig. 5 similar to that of Fig. 6, but illustrating another embodiment of the present disclosure.

DETAILED DESCRIPTION

For the purposes of promoting an understanding of the principles of the present disclosure, reference will now be made to the embodiments illustrated in the drawings, and specific language will be used to describe the same. It is nevertheless understood that no
5 limitation to the scope of the disclosure is intended. Any alterations and further modifications to the described devices, systems, and methods, and any further application of the principles of the present disclosure are fully contemplated and included within the present disclosure as would normally occur to one skilled in the art to which the disclosure relates. In particular, it is fully contemplated that the features, components, and/or steps described with
10 respect to one embodiment may be combined with the features, components, and/or steps described with respect to other embodiments of the present disclosure. For the sake of brevity, however, the numerous iterations of these combinations will not be described separately.

Referring to Figs. 1 and 2, shown therein is a vessel 100 having a stenosis according
15 to an embodiment of the present disclosure. In that regard, Fig. 1 is a diagrammatic perspective view of the vessel 100, while Fig. 2 is a partial cross-sectional perspective view of a portion of the vessel 100 taken along section line 2-2 of Fig. 1. Referring more specifically to Fig. 1, the vessel 100 includes a proximal portion 102 and a distal portion 104. A lumen 106 extends along the length of the vessel 100 between the proximal portion 102
20 and the distal portion 104. In that regard, the lumen 106 is configured to allow the flow of fluid through the vessel. In some instances, the vessel 100 is a blood vessel. In some particular instances, the vessel 100 is a coronary artery. In such instances, the lumen 106 is configured to facilitate the flow of blood through the vessel 100.

As shown, the vessel 100 includes a stenosis 108 between the proximal portion 102
25 and the distal portion 104. Stenosis 108 is generally representative of any blockage or other structural arrangement that results in a restriction to the flow of fluid through the lumen 106 of the vessel 100. Embodiments of the present disclosure are suitable for use in a wide variety of vascular applications, including without limitation coronary, peripheral (including but not limited to lower limb, carotid, and neurovascular), renal, and/or venous. Where the
30 vessel 100 is a blood vessel, the stenosis 108 may be a result of plaque buildup, including without limitation plaque components such as fibrous, fibro-lipidic (fibro fatty), necrotic core, calcified (dense calcium), blood, fresh thrombus, and mature thrombus. Generally, the composition of the stenosis will depend on the type of vessel being evaluated. In that regard,

it is understood that the concepts of the present disclosure are applicable to virtually any type of blockage or other narrowing of a vessel that results in decreased fluid flow.

Referring more particularly to Fig. 2, the lumen 106 of the vessel 100 has a diameter 110 proximal of the stenosis 108 and a diameter 112 distal of the stenosis. In some instances, the diameters 110 and 112 are substantially equal to one another. In that regard, the diameters 110 and 112 are intended to represent healthy portions, or at least healthier portions, of the lumen 106 in comparison to stenosis 108. Accordingly, these healthier portions of the lumen 106 are illustrated as having a substantially constant cylindrical profile and, as a result, the height or width of the lumen has been referred to as a diameter.

However, it is understood that in many instances these portions of the lumen 106 will also have plaque buildup, a non-symmetric profile, and/or other irregularities, but to a lesser extent than stenosis 108 and, therefore, will not have a cylindrical profile. In such instances, the diameters 110 and 112 are understood to be representative of a relative size or cross-sectional area of the lumen and do not imply a circular cross-sectional profile.

As shown in Fig. 2, stenosis 108 includes plaque buildup 114 that narrows the lumen 106 of the vessel 100. In some instances, the plaque buildup 114 does not have a uniform or symmetrical profile, making angiographic evaluation of such a stenosis unreliable. In the illustrated embodiment, the plaque buildup 114 includes an upper portion 116 and an opposing lower portion 118. In that regard, the lower portion 118 has an increased thickness relative to the upper portion 116 that results in a non-symmetrical and non-uniform profile relative to the portions of the lumen proximal and distal of the stenosis 108. As shown, the plaque buildup 114 decreases the available space for fluid to flow through the lumen 106. In particular, the cross-sectional area of the lumen 106 is decreased by the plaque buildup 114. At the narrowest point between the upper and lower portions 116, 118 the lumen 106 has a height 120, which is representative of a reduced size or cross-sectional area relative to the diameters 110 and 112 proximal and distal of the stenosis 108. Note that the stenosis 108, including plaque buildup 114 is exemplary in nature and should be considered limiting in any way. In that regard, it is understood that the stenosis 108 has other shapes and/or compositions that limit the flow of fluid through the lumen 106 in other instances. While the vessel 100 is illustrated in Figs. 1 and 2 as having a single stenosis 108 and the description of the embodiments below is primarily made in the context of a single stenosis, it is nevertheless understood that the devices, systems, and methods described herein have similar application for a vessel having multiple stenosis regions.

Referring now to Fig. 3, the vessel 100 is shown with instruments 130 and 132 positioned therein according to an embodiment of the present disclosure. In general, instruments 130 and 132 may be any form of device, instrument, or probe sized and shaped to be positioned within a vessel. In the illustrated embodiment, instrument 130 is generally representative of a guide wire, while instrument 132 is generally representative of a catheter. In that regard, instrument 130 extends through a central lumen of instrument 132. However, in other embodiments, the instruments 130 and 132 take other forms. In that regard, the instruments 130 and 132 are of similar form in some embodiments. For example, in some instances, both instruments 130 and 132 are guide wires. In other instances, both instruments 130 and 132 are catheters. On the other hand, the instruments 130 and 132 are of different form in some embodiments, such as the illustrated embodiment, where one of the instruments is a catheter and the other is a guide wire. Further, in some instances, the instruments 130 and 132 are disposed coaxial with one another, as shown in the illustrated embodiment of Fig. 3. In other instances, one of the instruments extends through an off-center lumen of the other instrument. In yet other instances, the instruments 130 and 132 extend side-by-side. In some particular embodiments, at least one of the instruments is as a rapid-exchange device, such as a rapid-exchange catheter. In such embodiments, the other instrument is a buddy wire or other device configured to facilitate the introduction and removal of the rapid-exchange device. Further still, in other instances, instead of two separate instruments 130 and 132 a single instrument is utilized. In that regard, the single instrument incorporates aspects of the functionalities (*e.g.*, data acquisition) of both instruments 130 and 132 in some embodiments.

Instrument 130 is configured to obtain diagnostic information about the vessel 100. In that regard, the instrument 130 includes one or more sensors, transducers, and/or other monitoring elements configured to obtain the diagnostic information about the vessel. The diagnostic information includes one or more of pressure, flow (velocity), images (including images obtained using ultrasound (*e.g.*, IVUS), OCT, thermal, and/or other imaging techniques), temperature, and/or combinations thereof. The one or more sensors, transducers, and/or other monitoring elements are positioned adjacent a distal portion of the instrument 130 in some instances. In that regard, the one or more sensors, transducers, and/or other monitoring elements are positioned less than 30 cm, less than 10 cm, less than 5 cm, less than 3 cm, less than 2 cm, and/or less than 1 cm from a distal tip 134 of the instrument 130 in some instances. In some instances, at least one of the one or more sensors, transducers, and/or other monitoring elements is positioned at the distal tip of the instrument 130.

The instrument 130 includes at least one element configured to monitor pressure within the vessel 100. The pressure monitoring element can take the form a piezo-resistive pressure sensor, a piezo-electric pressure sensor, a capacitive pressure sensor, an electromagnetic pressure sensor, a fluid column (the fluid column being in communication with a fluid column sensor that is separate from the instrument and/or positioned at a portion of the instrument proximal of the fluid column), an optical pressure sensor, and/or combinations thereof. In some instances, one or more features of the pressure monitoring element are implemented as a solid-state component manufactured using semiconductor and/or other suitable manufacturing techniques. Examples of commercially available guide wire products that include suitable pressure monitoring elements include, without limitation, the PrimeWire PRESTIGE® pressure guide wire, the PrimeWire® pressure guide wire, and the ComboWire® XT pressure and flow guide wire, each available from Volcano Corporation, as well as the PressureWire™ Certus guide wire and the PressureWire™ Aegis guide wire, each available from St. Jude Medical, Inc. Generally, the instrument 130 is sized such that it can be positioned through the stenosis 108 without significantly impacting fluid flow across the stenosis, which would impact the distal pressure reading. Accordingly, in some instances the instrument 130 has an outer diameter of 0.018" or less. In some embodiments, the instrument 130 has an outer diameter of 0.014" or less.

Instrument 132 is also configured to obtain diagnostic information about the vessel 100. In some instances, instrument 132 is configured to obtain the same diagnostic information as instrument 130. In other instances, instrument 132 is configured to obtain different diagnostic information than instrument 130, which may include additional diagnostic information, less diagnostic information, and/or alternative diagnostic information. The diagnostic information obtained by instrument 132 includes one or more of pressure, flow (velocity), images (including images obtained using ultrasound (*e.g.*, IVUS), OCT, thermal, and/or other imaging techniques), temperature, and/or combinations thereof. Instrument 132 includes one or more sensors, transducers, and/or other monitoring elements configured to obtain this diagnostic information. In that regard, the one or more sensors, transducers, and/or other monitoring elements are positioned adjacent a distal portion of the instrument 132 in some instances. In that regard, the one or more sensors, transducers, and/or other monitoring elements are positioned less than 30 cm, less than 10 cm, less than 5 cm, less than 3 cm, less than 2 cm, and/or less than 1 cm from a distal tip 136 of the instrument 132 in some instances. In some instances, at least one of the one or more sensors,

transducers, and/or other monitoring elements is positioned at the distal tip of the instrument 132.

Similar to instrument 130, instrument 132 also includes at least one element configured to monitor pressure within the vessel 100. The pressure monitoring element can take the form a piezo-resistive pressure sensor, a piezo-electric pressure sensor, a capacitive pressure sensor, an electromagnetic pressure sensor, a fluid column (the fluid column being in communication with a fluid column sensor that is separate from the instrument and/or positioned at a portion of the instrument proximal of the fluid column), an optical pressure sensor, and/or combinations thereof. In some instances, one or more features of the pressure monitoring element are implemented as a solid-state component manufactured using semiconductor and/or other suitable manufacturing techniques. Currently available catheter products suitable for use with one or more of Siemens AXIOM Sensis, Mennen Horizon XVu, and Philips Xper IM Physiomonitoring 5 and that include pressure monitoring elements can be utilized for instrument 132 in some instances.

In accordance with aspects of the present disclosure, at least one of the instruments 130 and 132 is configured to monitor a pressure within the vessel 100 distal of the stenosis 108 and at least one of the instruments 130 and 132 is configured to monitor a pressure within the vessel proximal of the stenosis. In that regard, the instruments 130, 132 are sized and shaped to allow positioning of the at least one element configured to monitor pressure within the vessel 100 to be positioned proximal and/or distal of the stenosis 108 as necessary based on the configuration of the devices. In that regard, Fig. 3 illustrates a position 138 suitable for measuring pressure distal of the stenosis 108. The position 138 is less than 5 cm, less than 3 cm, less than 2 cm, less than 1 cm, less than 5 mm, and/or less than 2.5 mm from the distal end of the stenosis 108 (as shown in Fig. 2) in some instances. Fig. 3 also illustrates a plurality of suitable positions for measuring pressure proximal of the stenosis 108. In that regard, positions 140, 142, 144, 146, and 148 each represent a position that is suitable for monitoring the pressure proximal of the stenosis in some instances. In that regard, the positions 140, 142, 144, 146, and 148 are positioned at varying distances from the proximal end of the stenosis 108 ranging from more than 20 cm down to about 5 mm or less. Generally, the proximal pressure measurement will be spaced from the proximal end of the stenosis. Accordingly, in some instances, the proximal pressure measurement is taken at a distance equal to or greater than an inner diameter of the lumen of the vessel from the proximal end of the stenosis. In the context of coronary artery pressure measurements, the proximal pressure measurement is generally taken at a position proximal of the stenosis and

distal of the aorta, within a proximal portion of the vessel. However, in some particular instances of coronary artery pressure measurements, the proximal pressure measurement is taken from a location inside the aorta. In other instances, the proximal pressure measurement is taken at the root or ostium of the coronary artery. In some instances, the proximal pressure measurement is referred to as the aortic pressure.

Referring now to Fig. 4, shown therein is a system 150 according to an embodiment of the present disclosure. In that regard, Fig. 4 is a diagrammatic, schematic view of the system 150. As shown, the system 150 includes an instrument 152. In that regard, in some instances instrument 152 is suitable for use as at least one of instruments 130 and 132 discussed above. Accordingly, in some instances the instrument 152 includes features similar to those discussed above with respect to instruments 130 and 132 in some instances. In the illustrated embodiment, the instrument 152 is a guide wire having a distal portion 154 and a housing 156 positioned adjacent the distal portion. In that regard, the housing 156 is spaced approximately 3 cm from a distal tip of the instrument 152. The housing 156 is configured to house one or more sensors, transducers, and/or other monitoring elements configured to obtain the diagnostic information about the vessel. In the illustrated embodiment, the housing 156 contains at least a pressure sensor configured to monitor a pressure within a lumen in which the instrument 152 is positioned. A shaft 158 extends proximally from the housing 156. A torque device 160 is positioned over and coupled to a proximal portion of the shaft 158. A proximal end portion 162 of the instrument 152 is coupled to a connector 164. A cable 166 extends from connector 164 to a connector 168. In some instances, connector 168 is configured to be plugged into an interface 170. In that regard, interface 170 is a patient interface module (PIM) in some instances. In some instances, the cable 166 is replaced with a wireless connection. In that regard, it is understood that various communication pathways between the instrument 152 and the interface 170 may be utilized, including physical connections (including electrical, optical, and/or fluid connections), wireless connections, and/or combinations thereof.

The interface 170 is communicatively coupled to a computing device 172 via a connection 174. Together, connector 164, cable 166, connector 168, interface 170, and connection 174 facilitate communication between the one or more sensors, transducers, and/or other monitoring elements of the instrument 152 and the computing device 172. However, this communication pathway is exemplary in nature and should not be considered limiting in any way. In that regard, it is understood that any communication pathway between the instrument 152 and the interface 170 may be utilized, including physical

connections (including electrical, optical, and/or fluid connections), wireless connections, and/or combinations thereof. Similarly, it is understood that any communication pathway between the interface 170 and the computing device 172 may be utilized, including physical connections (including electrical, optical, and/or fluid connections), wireless connections, and/or combinations thereof. Accordingly, it is understood that additional components (*e.g.*, connectors, antennas, routers, switches, etc.) not illustrated in Fig. 4 may be included to facilitate communication between the instrument 152, the interface 170, and the computing device 172.

In some embodiments, the connection 174 is a wireless connection. In some instances, the connection 174 includes a communication link over a network (*e.g.*, intranet, internet, telecommunications network, and/or other network). In that regard, it is understood that the computing device 172 is positioned remote from an operating area where the instrument 152 is being used in some instances. Having the connection 174 include a connection over a network can facilitate communication between the instrument 152 and the remote computing device 172 regardless of whether the computing device is in an adjacent room, an adjacent building, or in a different state/country. Further, it is understood that the communication pathway between the instrument 152 and the computing device 172 is a secure connection in some instances. Further still, it is understood that, in some instances, the data communicated over one or more portions of the communication pathway between the instrument 152 and the computing device 172 is encrypted.

The system 150 also includes an instrument 175. In that regard, in some instances instrument 175 is suitable for use as at least one of instruments 130 and 132 discussed above. Accordingly, in some instances the instrument 175 includes features similar to those discussed above with respect to instruments 130 and 132. In the illustrated embodiment, the instrument 175 is a catheter-type device. In that regard, the instrument 175 includes one or more sensors, transducers, and/or other monitoring elements adjacent a distal portion of the instrument configured to obtain the diagnostic information about the vessel. In the illustrated embodiment, the instrument 175 includes a pressure sensor configured to monitor a pressure within a lumen in which the instrument 175 is positioned. In one particular embodiment, instrument 175 is a pressure-sensing catheter that includes a fluid column extending along its length. In such an embodiment, a hemostasis valve is fluidly coupled to the fluid column of the catheter, a manifold is fluidly coupled to the hemostasis valve, and tubing extends between the components as necessary to fluidly couple the components. In that regard, the fluid column of the catheter is in fluid communication with a pressure sensor via the valve,

manifold, and tubing. In some instances, the pressure sensor is part of or in communication with computing device 172. In other instances, the pressure sensor is a separate component positioned between the instrument 175 and the interface 170 or between the interface 170 and the computing device 172. In some instances, the computing device 172 is a hemodynamic
5 system such as Siemens AXIOM Sensis, Mennen Horizon XVu, and Philips Xper IM Physiomonitoring 5. The instrument 175 is in communication with the interface 170 via connection 177. The interface 170, in turn, is communicatively coupled to the computing device 172 via a connection 178.

Similar to the connections between instrument 152 and the interface 170 and the
10 computing device 172, connections 177 and 178 facilitate communication between the one or more sensors, transducers, and/or other monitoring elements of the instrument 175 and the interface 170 and the computing device 172. Again, however, this communication pathway is exemplary in nature and should not be considered limiting in any way. In that regard, it is understood that any communication pathway between the instrument 175 and the interface
15 170 may be utilized, including physical connections (including electrical, optical, and/or fluid connections), wireless connections, and/or combinations thereof. Similarly, it is understood that any communication pathway between the interface 170 and the computing device 172 may be utilized, including physical connections (including electrical, optical, and/or fluid connections), wireless connections, and/or combinations thereof. Accordingly, it is
20 understood that additional components (*e.g.*, connectors, antennas, routers, switches, etc.) not illustrated in Fig. 4 may be included to facilitate communication between the instrument 175, the interface 170, and the computing device 172.

In some embodiments, the connection 178 is a wireless connection. In some instances, the connection 178 includes a communication link over a network (*e.g.*, intranet,
25 internet, telecommunications network, and/or other network). In that regard, it is understood that the computing device 172 is positioned remote from an operating area where the instrument 175 is being used in some instances. Having the connection 178 include a connection over a network can facilitate communication between the instrument 175 and the remote computing device 172 regardless of whether the computing device is in an adjacent
30 room, an adjacent building, or in a different state/country. Further, it is understood that the communication pathway between the instrument 175 and the computing device 172 is a secure connection in some instances. Further still, it is understood that, in some instances, the data communicated over one or more portions of the communication pathway between the instrument 175 and the computing device 172 is encrypted.

It is understood that one or more components of the system 150 are not included, are implemented in a different arrangement/order, and/or are replaced with an alternative device/mechanism in other embodiments of the present disclosure. Alternatively, additional components and/or devices may be implemented into the system. Generally speaking, the communication pathway between either or both of the instruments 152, 175 and the computing device 172 may have no intermediate nodes (*i.e.*, a direct connection), one intermediate node between the instrument and the computing device, or a plurality of intermediate nodes between the instrument and the computing device.

In some embodiments, the interface 170 includes a processor and random access memory and is programmed to execute steps associated with the data acquisition and analysis described herein. In particular, in some embodiments the interface 170 is configured to receive and display pressure readings from one or both of the instruments 152 and 175 and/or calculate (and display) FFR or other pressure differential based on the pressure measurements obtained from the instruments 152 and 175. Accordingly, it is understood that any steps related to data acquisition, data processing, instrument control, and/or other processing or control aspects of the present disclosure, including those incorporated by reference, may be implemented by the interface 170 using corresponding instructions stored on or in a non-transitory computer readable medium accessible by the computing device. In some embodiments, the interface 170 includes one or more processing and/or signal conditioning features and/or associated components/circuitry as described in U.S. Patent No. 6,585,660, which is hereby incorporated by reference in its entirety.

In the illustrated embodiment of Fig. 4, the interface 170 includes a housing 180. The housing 180 contains the electronic components of the interface 170. In that regard, exemplary embodiments of electronic component arrangements suitable for interface 170 as described below with respect to Figs. 5 and 6. In some embodiments, the interface 170 is sized to be handheld and/or sized to be positioned on or near a patient bed (*e.g.*, attached to a bed rail or IV pole). In that regard, in some instances the interface 170 is similar in size to the SmartMap® Pressure Instrument available from Volcano Corporation, which has housing dimensions of approximately 15.75 cm (6.3”) wide, 8.853 cm (3.54”) tall, and 4.48 cm (1.79”) deep. Generally, the interface 170 has a width between about 5 cm and about 25 cm, a height between about 5 cm and about 25 cm, and a depth between about 1 cm and about 10 cm.

The interface 170 also includes a display 182 and buttons 184. In that regard, the display 182 is configured to display various diagnostic information such as distal pressure,

proximal pressure, pressure differentials (including FFR), distal pressure waveforms, proximal pressure waveforms, and/or additional diagnostic parameters. In some embodiments, in order to conserve the amount of energy needed for operation of the interface 170, the display 182 is a low-power display, such as a liquid crystal display. However, any type of visual display may be utilized, including color and/or monochromatic displays. In some instances, the display 182 covers a majority of a front surface of the display. In that regard, in some particular embodiments the display 182 is a touchscreen. In such embodiments, the buttons 184 may be virtual buttons (*i.e.*, displayed on the touchscreen display 182), physical buttons, and/or combinations thereof. Generally, the buttons 184 are configured to facilitate configuration and operation of the interface 170. It is understood that any number of buttons may be utilized and that buttons may be utilized for multiple functionalities and/or be dedicated to a single function. As a result, the interface 170 may include one or more virtual or physical buttons configured to facilitate use of the interface in the manners described herein.

Referring now to Fig. 5, shown therein is a schematic of the interface 170 according to an exemplary embodiment of the present disclosure. In that regard, the interface 170 includes an input connector 190 for receiving signals from a distal pressure sensing component 191. Accordingly, in some embodiments with an arrangement similar to that shown in Fig. 4, input connector 190 is configured to receive the connector 168 that is communication with instrument 152, where distal pressure sensing component 191 is a pressure sensing component of the instrument 152. The interface 170 also includes an output connector 192 configured to send a distal pressure signal to a distal pressure input 193 of a hemo system or other computing device. Accordingly, in some embodiments with an arrangement similar to that shown in Fig. 4, output connector 192 is configured to send the distal pressure signal to an input of computing device 172 over connection 174. In that regard, in some embodiments the distal pressure signal is modulated based on the hemo system's excitation voltage to provide a low level output of the distal pressure signal to the hemo system. A low level output in this context is typically $5\mu\text{V}/V_{\text{exc}}/\text{mmHg}$, where V_{exc} is the excitation voltage. However, larger or smaller level outputs are used in some instances.

In some embodiments, the output connector 192 is also used to facilitate energy harvesting from the hemo system or other pressure measuring system. In that regard, to eliminate the need for an additional power supply within the interface 170, a power supply circuit 220 extracts a small amount of power from the hemo system's excitation voltage associated with the distal pressure input 193. The power supply circuit 220 converts the

extracted energy into the power needed to run the remaining circuitry of the interface 170. In some instances, the power supply circuit 220 is configured to be the only power source used to power the components of interface 170. Since the excitation signal can be AC, positive or negative DC, and/or have various wave form shapes and voltages, the power supply circuit
5 220 must be able to accept these and convert to a regulated power supply. In that regard, the voltage extracted from the excitation signal is converted to a regulated Vcc voltage to operate the low power circuitry using a buck or boost regulator depending on the input voltage. A current limiter minimizes distortion to the hemo system's waveform at the peaks. In some instances, the current is limited to a level below the AAMI transducer limits as to be
10 compatible with most hemo systems. In some instances, the hemo system's excitation voltage meets the IEC 60601-2-34 standard. In some alternative embodiments, the power supply circuit is configured to interface with a battery or other rechargeable power supply device that can be utilized to power the components of the interface. In some alternative embodiments, the power supply circuit is configured to interface with an AC adapter that is to
15 be plugged into a wall outlet in order to provide power to the components of the interface.

The interface 170 also includes an input/output connectors 194 and 195 for interfacing with a proximal pressure measurement system. In some particular embodiments, the input/output connectors 194 and 195 are configured to work with a pressure monitoring device of a hemo stat system. Generally, the input/output connector 194 is configured to
20 receive signals from a proximal pressure sensing component 196. Accordingly, in some embodiments with an arrangement similar to that shown in Fig. 4, input/output connector 194 is configured to receive signals from instrument 175, where the proximal pressure sensing component 196 is a pressure sensing component associated with the instrument 175. The input/output connector 195 is configured to send a proximal pressure signal to a proximal
25 pressure input 197 of a hemo system or other computing device. Accordingly, in some embodiments with an arrangement similar to that shown in Fig. 4, the input/output connector 195 is configured to send the proximal pressure signal to an input of computing device 172 over connection 178.

In the illustrated embodiment of Fig. 5, conductors 200 and 202 carry the excitation
30 signal to the proximal pressure sensing component 196. An amplifier 204 is electrically connected to the conductors 200 and 202 as shown. The amplifier 204 is an operational amplifier in some embodiments. The excitation signal extracted by amplifier 204 is sent to a microprocessor 206. As will be discussed below, the excitation signal is utilized to evaluate the proximal pressure signals received from the proximal pressure sensing component 196.

In the illustrated embodiment of Fig. 5, conductors 208 and 210 carry the proximal pressure signal from the proximal pressure sensing component 196 back to the proximal pressure input 197 of the computing device. In that regard, an amplifier 212 is electrically connected to the conductors 208 and 210 as shown. The amplifier 212 is an operational
5 amplifier in some embodiments. The amplifier 212 is configured to monitor or sample the proximal pressure signal being supplied from the proximal pressure sensing component 196. The sampled proximal pressure signal is then sent to the microprocessor 206. Accordingly, both the excitation signal/voltage sampled from conductors 200 and 202 and the proximal pressure signal sampled from conductors 208 and 210 are fed to the microprocessor 206. In
10 some instances, the microprocessor 206 calculates the proximal pressure based on the excitation signal voltage (V_{exc}), and the proximal pressure sensing component's output. In that regard, the proximal pressure sensing component's output conforms to the AAMI standard of $5\mu V/V_{exc}/\text{mmHg}$ in some instances. For an AC excitation signal, the microprocessor 206 must measure the proximal pressure signal voltage in synchrony with the
15 excitation waveform. In some instances, rather than the low-level inputs described above, the proximal pressure signal is received by the interface 170 as a high level signal. For example, the proximal pressure signal is a high level signal from a Volcano LoMap (available from Volcano Corporation) or from an external hemo system.

In some embodiments, the input/output connectors 194 and 195 are also used to
20 facilitate energy harvesting from the hemo system or other pressure measuring system. In that regard, to eliminate the need for an additional power supply within the interface 170, the power supply circuit 220 may be connected to conductors 200 and 202 and utilized to extract a small amount of power from the hemo system's excitation voltage for the proximal pressure sensing component 196 and convert it into the power needed to run the remaining circuitry of
25 the interface 170. As noted above, when connected to the proximal pressure sensing side the power supply circuit 220 is still configured to extract power from the excitation signal sent from the controller/computing device such that the extracted power can be used to power the components of interface 170. Since the excitation signal can be AC, positive or negative DC, and/or have various wave form shapes and voltages, the power supply circuit 220 must be
30 able to accept these and convert to a regulated power supply without distorting the waveform that continues on to the proximal pressure sensing component 196. This is necessary to avoid affecting the pressure measurements obtained by the proximal pressure sensing component 196. The voltage extracted from the excitation signal is converted to a regulated V_{cc} voltage

to operate the low power circuitry using a buck or boost regulator depending on the input voltage.

As noted above, the interface 170 is also configured to receive and process distal pressure signals from a distal pressure sensing component 191. In that regard, a signal conditioning portion 216 of the interface 170 is in communication with input 190 that receives the distal pressure signal. Referring now to Fig. 6, shown therein is a schematic of a portion the interface 170 according to an exemplary embodiment of the present disclosure. In particular, Fig. 6 shows a schematic of an exemplary embodiment of signal conditioning portion 216 of the interface 170. In that regard, the signal condition portion 216 is configured to condition signals received from the distal pressure measurement device. The signal conditioning portion 216 provides the excitation and amplification required for the distal pressure measurement device's pressure sensors, Ra and Rb, which collective form distal pressure sensing component 191 in some instances.

Calibration coefficients provided by the distal pressure measurement device utilizing an EPROM in the device connector, for example, are read to adjust the gain, offset, and temperature sensitivity for the device. The read values are used to adjust the three Digital to Analog Converters (DACs) 224, 226, and 228, in the distal pressure front end circuitry 216 that control the gain, offset, and temperature (TC) compensation, respectively. The distal pressure signal is then digitized with an Analog to Digital Converter 230, ADC, and sent to the microprocessor 206. The microprocessor 206 can then display the distal pressure, display a waveform of the distal pressure, or utilize the distal pressure or the distal pressure waveform for additional calculations. For example, in some instances the microprocessor utilizes the distal pressure and/or distal pressure waveform with the proximal pressure and/or proximal pressure waveform to calculate FFR, calculate a pressure differential between the proximal and distal pressures, identify a suitable diagnostic window for performing a pressure differential calculation without administering a hyperemic agent to the patient, calculate a pressure differential during the identified diagnostic window, and/or combinations thereof.

Referring now to Fig. 7, shown therein is a schematic of a portion the interface 170 according to another exemplary embodiment of the present disclosure. In particular, Fig. 7 shows a schematic of an exemplary embodiment of signal conditioning portion 216' of the interface 170. In that regard, the signal condition portion 216' is configured to condition signals received from the distal pressure measurement device. The signal conditioning portion 216' provides the excitation and amplification required for the distal pressure

measurement device's pressure sensors, Ra and Rb, which collectively form distal pressure sensing component 191 in some instances. The distal pressure signal from the pressure sensors, Ra and Rb, is digitized with a two-channel Analog to Digital Converter 230, ADC, and sent to the microprocessor 206 for the gain, offset, and/or temperature compensation.

5 Calibration coefficients provided by the distal pressure measurement device utilizing an EPROM 222 in the device connector, for example, are read to adjust the gain, offset, and/or temperature sensitivity for the device. The read values are used by the microprocessor 206 to control the gain, offset, and/or temperature compensation. Firmware within the microprocessor is utilized to control these parameters in some instances. The microprocessor
10 206 can then display the distal pressure, display a waveform of the distal pressure, or utilize the distal pressure or the distal pressure waveform for additional calculations. For example, in some instances the microprocessor utilizes the distal pressure and/or distal pressure waveform with the proximal pressure and/or proximal pressure waveform to calculate FFR, calculate a pressure differential between the proximal and distal pressures, identify a suitable
15 diagnostic window for performing a pressure differential calculation without administering a hyperemic agent to the patient, calculate a pressure differential during the identified diagnostic window, and/or combinations thereof. Accordingly, the signal conditioning portion 216' of Fig. 7 provides similar functionality to the signal conditioning device 216 of Fig. 6, but without the need for the three Digital to Analog Converters (DACs) 224, 226, and
20 228.

Referring again to Fig. 5, the interface 170 is also configured to output the distal pressure signal to an input 193 of a computing device. In that regard, the microprocessor 206 provides a digitized signal to an additional set of DACs in the distal output circuitry 218 that modulate the excitation of the hemo system to provide a proportional distal waveform of the
25 distal pressure voltage back to the hemo system through output 192. In some embodiments, the scaled voltage returned is the same as a standard proximal pressure transducer, $5\mu\text{V}/V_{\text{exc}}/\text{mmHg}$, per the AAMI standards. The output stage 218 modulates the external excitation of the hemo system to provide a duplicate wave shape, or a DC voltage, scaled to $5\mu\text{V} / V_{\text{exc}} / \text{mmHg}$, per AAMI standards for aortic transducers of the distal pressure for the
30 hemo system. Accordingly, by outputting the distal pressure signal through output 192 and the proximal pressure signal through output 196, both proximal and distal pressures can then be observed on the hemo system's display using the hemo system's standard low-level inputs.

As noted above, the interface 170 uses the proximal and distal pressure data received from the instruments to calculate and display information that can be useful in the evaluation

of the vessel and, in particular, evaluation of a stenosis of the vessel. In some instances, the interface is configured to calculate and display FFR. For an FFR measurement, the microprocessor first normalizes the distal pressure to the aortic pressure. The distal and aortic pressures will become disparaging as the distal pressure wire crosses the arterial lesion.

5 The peak difference between the two pressures is captured automatically, or with a manual button press, and an FFR calculation started. The resultant number is shown on the display. The peak difference is typically measured during hyperemia with the use of drugs like adenosine

In some embodiments the interface 170 includes external user-controlled buttons, such as buttons 184 shown in Fig. 4. In one particular embodiment, one of the buttons causes the microprocessor to 'normalize' the distal pressure measurement to the proximal pressure measurement. This is typically performed with the pressure sensing components 191 and 196 positioned in close proximity to one another within the patient such that they are subjected to similar pressures. In some instances, this calibration is performed proximal of the lesion and before the distal pressure sensing component 191 is advanced distally beyond the lesion. After the distal pressure sensing component 191 is placed beyond the suspect lesion actuation of another button causes the microprocessor 206 to calculate the ratio of the distal pressure to the proximal pressure, which provides an FFR value or pressure differential. In that regard, in some implementations the button is pressed by a user at a precise moment during hyperemia based on observation of the proximal and distal waveforms, which may be displayed on a separate device (*e.g.*, a display of the hemo system) or displayed on display 182 of interface 170. Alternatively, the determination of the appropriate moment for the FFR calculation can be done automatically by the microprocessor 206. In that regard, in some instances the FFR calculation is performed at a point coinciding with the peak difference between the distal and proximal (aortic) pressures. In some embodiments, a pressure differential is calculated during a diagnostic window without application of a hyperemic agent, as discussed below. In such embodiments, the pressure measurements and/or the pressure differential may be displayed continuously.

In that regard, in some instances the interface 170 is configured to provide pressure measurements and/or pressure differentials based on evaluation techniques as described in one or more of UK Patent Application Publication No. GB 2479340 A, filed March 10, 2010 and titled "METHOD AND APPARATUS FOR THE MEASUREMENT OF A FLUID FLOW RESTRICTION IN A VESSEL", UK Patent Application No. 1100137.7 filed January 6, 2011 and titled "APPARATUS AND METHOD OF ASSESSING A NARROWING IN A

FLUID FILLED TUBE”, U.S. Provisional Patent Application No. 61/525,739, filed on August 20, 2011 and titled “DEVICES, SYSTEMS AND METHODS FOR ASSESSING A VESSEL,” and U.S. Provisional Patent Application No. 61/525,736, filed on August 20, 2011 and titled “DEVICES, SYSTEMS, AND METHODS FOR VISUALLY DEPICTING A VESSEL AND EVALUATING TREATMENT OPTIONS,” each of which is hereby incorporated by reference in its entirety.

In some embodiments, the interface 170 is utilized to calculate and display FFR in a traditional FFR procedure where the patient is administered a hyperemic agent. In other embodiments, the interface 170 is utilized to calculate a pressure differential similar to FFR (*i.e.*, the ratio of distal pressure to proximal pressure) but without the use of a hyperemic agent. In that regard, a suitable diagnostic window for making such calculations must be determined and/or identified to have a useful measurement. The diagnostic window for evaluating differential pressure across a stenosis without the use of a hyperemic agent in accordance with the present disclosure may be identified based on characteristics and/or components of one or more of proximal pressure measurements, distal pressure measurements, proximal velocity measurements, distal velocity measurements, ECG waveforms, and/or other identifiable and/or measurable aspects of vessel performance. In that regard, various signal processing and/or computational techniques can be applied to the characteristics and/or components of one or more of proximal pressure measurements, distal pressure measurements, proximal velocity measurements, distal velocity measurements, ECG waveforms, and/or other identifiable and/or measurable aspects of vessel performance to identify a suitable diagnostic window.

In some embodiments, the determination of the diagnostic window and/or the calculation of the pressure differential are performed in approximately real time or live to identify the diagnostic window and calculate the pressure differential. In that regard, calculating the pressure differential in “real time” or “live” within the context of the present disclosure is understood to encompass calculations that occur within 10 seconds of data acquisition. It is recognized, however, that often “real time” or “live” calculations are performed within 1 second of data acquisition. In some instances, the “real time” or “live” calculations are performed concurrent with data acquisition. In some instances the calculations are performed by a processor in the delays between data acquisitions. For example, if data is acquired from the pressure sensing devices for 1 ms every 5 ms, then in the 4 ms between data acquisitions the processor can perform the calculations. It is understood that these timings are for example only and that data acquisition rates, processing

times, and/or other parameters surrounding the calculations will vary. In other embodiments, the pressure differential calculation is performed 10 or more seconds after data acquisition. For example, in some embodiments, the data utilized to identify the diagnostic window and/or calculate the pressure differential are stored for later analysis.

5 In some instances, the diagnostic window is selected by identifying a portion of the cardiac cycle corresponding to a time period in which the change in velocity (*i.e.*, dU) fluctuates around zero. Time periods where the change in velocity is relatively constant and approximately zero (*i.e.*, the speed of the fluid flow is stabilized) are suitable diagnostic windows for evaluating a pressure differential across a stenosis of a vessel without the use of
10 a hyperemic agent in accordance with the present disclosure. In that regard, in a fluid flow system, the separated forward and backward generated pressures are defined by:

$$dP_+ = \frac{1}{2}(dP + \rho c dU) \text{ and } dP_- = \frac{1}{2}(dP - \rho c dU),$$

where dP is the differential of pressure, ρ is the density of the fluid within the vessel, c is the wave speed, and dU is the differential of flow velocity. However, where the flow velocity of
15 the fluid is substantially constant, dU is approximately zero and the separated forward and backward generated pressures are defined by:

$$dP_+ = \frac{1}{2}(dP + \rho c(0)) = \frac{1}{2}dP \text{ and } dP_- = \frac{1}{2}(dP - \rho c(0)) = \frac{1}{2}dP.$$

In other words, during the time periods where dU is approximately zero, the forward and backward generated pressures are defined solely by changes in pressure.

20 Accordingly, during such time periods the severity of a stenosis within the vessel can be evaluated based on pressure measurements taken proximal and distal of the stenosis. In that regard, by comparing the forward and/or backward generated pressure distal of a stenosis to the forward and/or backward generated pressure proximal of the stenosis, an evaluation of the severity of the stenosis can be made. For example, the forward-generated pressure

25 differential can be calculated as $\frac{dP_{+distal}}{dP_{+proximal}}$, while the backward-generated pressure

differential can be calculated as $\frac{dP_{-distal}}{dP_{-proximal}}$.

In the context of the coronary arteries, a forward-generated pressure differential is utilized to evaluate a stenosis in some instances. In that regard, the forward-generated pressure differential is calculated based on proximally originating (*i.e.*, originating from the
30 aorta) separated forward pressure waves and/or reflections of the proximally originating

separated forward pressure waves from vascular structures distal of the aorta in some instances. In other instances, a backward-generated pressure differential is utilized in the context of the coronary arteries to evaluate a stenosis. In that regard, the backward-generated pressure differential is calculated based on distally originating (*i.e.*, originating from the
5 microvasculature) separated backward pressure waves and/or reflections of the distally originating separated backward pressure waves from vascular structures proximal of the microvasculature.

In yet other instances, a pressure wave is introduced into the vessel by an instrument or medical device. In that regard, the instrument or medical device is utilized to generate a
10 proximally originating forward pressure wave, a distally originating backward pressure wave, and/or combinations thereof for use in evaluating the severity of the stenosis. For example, in some embodiments an instrument having a movable membrane is positioned within the vessel. The movable membrane of the instrument is then activated to cause movement of the membrane and generation of a corresponding pressure wave within the fluid of the vessel.
15 Based on the configuration of the instrument, position of the membrane within the vessel, and/or the orientation of the membrane within the vessel the generated pressure wave(s) will be directed distally, proximally, and/or both. Pressure measurements based on the generated pressure wave(s) can then be analyzed to determine the severity of the stenosis.

There are a variety of signal processing techniques that can be utilized to identify time
20 periods where the change in velocity is relatively constant and approximately zero, including using a differential, first derivative, second derivative, and/or third derivative of the velocity measurement are utilized. For example, identifying time periods during the cardiac cycle where the first derivative of velocity is relatively constant and approximately zero allows the localization of time periods where velocity is relatively constant. Further, identifying time
25 periods during the cardiac cycle where the second derivative of velocity is relatively constant and approximately zero allows the localization of a time period where acceleration is relatively constant and near zero, but not necessarily zero.

While examples of specific techniques for selecting a suitable diagnostic window have been described above, it is understood that these are exemplary and that other
30 techniques may be utilized. In that regard, it is understood that the diagnostic window is determined using one or more techniques selected from: identifying a feature of a waveform or other data feature and selecting a starting point relative to the identified feature (*e.g.*, before, after, or simultaneous with the feature); identifying a feature of a waveform or other data feature and selecting an ending point relative to the identified feature (*e.g.*, before, after,

or simultaneous with the feature); identifying a feature of a waveform or other data feature and selecting a starting point and an ending point relative to the identified feature; identifying a starting point and identifying an ending point based on the starting point; and identifying an ending point and indentifying a starting point based on the ending point. Additional details of techniques for selecting a suitable diagnostic window are described in U.S. Provisional Patent Application No. 61/525,739, filed on August 20, 2011 and titled “DEVICES, SYSTEMS AND METHODS FOR ASSESSING A VESSEL,” which is hereby incorporated by reference in its entirety. In that regard, it is understood that the interface 170 may be programmed to determine one or more diagnostic windows based on the techniques described in the present disclosure, including those incorporated by reference, and/or include one or more hardware features configured to identify one or more diagnostic windows based on the techniques described in the present disclosure, including those incorporated by reference.

Further, for a variety of reasons the proximal pressure measurements and the distal pressure measurements received by the interface 170 are not temporally aligned in some instances. For example, during data acquisition, there will often be a delay between the distal pressure measurement signals and the proximal pressure measurement signals due to hardware signal handling differences between the instrument(s) utilized to obtain the measurements. In that regard, the differences can come from physical sources (such as cable length and/or varying electronics) and/or can be due to signal processing differences (such as filtering techniques). The resulting delay between the signals is between about 5 ms and about 150 ms in some instances. Because individual cardiac cycles may last between about 500 ms and about 1000 ms and the diagnostic window may be a small percentage of the total length of the cardiac cycle, longer delays between the proximal and distal pressure measurement signals can have a significant impact on alignment of the pressure data for calculating a pressure differential for a desired diastolic window of a cardiac cycle.

As a result, in some instances, it is necessary to shift one of the proximal and distal pressures relative to the other of the distal and proximal pressures in order to temporally align the pressure measurements. For example, a portion of the distal pressure measurement or proximal pressure measurement may be shifted to be temporally aligned with a corresponding portion of the proximal pressure measurement or distal pressure measurement, respectively, coinciding with the diagnostic window. While a shift of only a portion of the distal or proximal pressure measurement associated with the diagnostic window is utilized in some instances, in other instances all or substantially all of the proximal and distal pressures are aligned before the portions corresponding to a selected diagnostic window are identified.

Alignment of all or portion(s) of the proximal and distal pressures is accomplished using a hardware approach in some instances. For example, one or more hardware components are positioned within the communication path of the proximal pressure measurement, the distal pressure measurement, and/or both to provide any necessary delays to temporally align the received pressure signals. In some instances, these hardware components are positioned within the interface 170. In other instances, alignment of all or portion(s) of the proximal and distal pressures is accomplished using a software approach. For example, a cross-correlation function or matching technique is utilized to align the cardiac cycles in some embodiments. In other embodiments, the alignment is based on a particular identifiable feature of the cardiac cycle, such as an ECG R-wave or a pressure peak. Additionally, in some embodiments alignment is performed by a software user where adjustments are made to the delay time of at least one of the proximal and distal pressures until the cardiac cycles are visually aligned to the user. A further technique for aligning the signals is to apply a synchronized timestamp at the point of signal acquisition. Further, in some instances combinations of one or more of hardware, software, user, and/or time-stamping approaches are utilized to align the signals.

Regardless of the manner of implementation, several approaches are available for the aligning the proximal and distal pressure measurement signals. In some instances, each individual distal pressure measurement cardiac cycle is individually shifted to match the corresponding proximal pressure measurement cardiac cycle. In other instances, an average shift for a particular procedure is calculated at the beginning of the procedure and all subsequent cardiac cycles during the procedure are shifted by that amount. This technique requires little processing power for implementation after the initial shift is determined, but can still provide a relatively accurate alignment of the signals over the course of a procedure because the majority of the signal delay is due to fixed sources that do not change from patient to patient or within the procedure. In yet other instances, a new average shift is calculated each time that the proximal and distal pressure signals are normalized to one another during a procedure. In that regard, one or more times during a procedure the sensing element utilized for monitoring pressure distal of the stenosis is positioned adjacent the sensing element utilized for monitoring pressure proximal of the stenosis such that both sensing elements should have the same pressure reading. If there is a difference between the pressure readings, then the proximal and distal pressure signals are normalized to one another. As a result, the subsequently obtained proximal and distal pressure measurements

are more consistent with each other and, therefore, the resulting pressure differential calculations are more accurate.

With the proximal and distal pressure measurements aligned, the pressure differential for the diagnostic window is calculated. In some instances, the pressure differential is
5 calculated using average values for the proximal and distal pressure measurements across the diagnostic window. The pressure differential calculations of the present disclosure are performed for a single cardiac cycle, in some instances. In other instances, the pressure differential calculations are performed for multiple cardiac cycles. In that regard, accuracy of the pressure differential can be improved by performing the pressure differential calculations
10 over multiple cardiac cycles and averaging the values and/or using an analysis technique to identify one or more of the calculated values that is believed to be most and/or least accurate.

One advantage of the techniques of the present disclosure for identifying diagnostic windows and evaluating pressure differentials is the concept of “beat matching”. In that regard, the proximal and distal waveforms for the same cardiac cycle are analyzed together
15 with no averaging or individual calculations that span more than a single cardiac cycle. As a result, interruptions in the cardiac cycle (such as ectopic heartbeats) equally affect the proximal and distal recordings. As a result, these interruptions that can be detrimental to current FFR techniques have minor effect on the techniques of the present disclosure. Further, in some embodiments of the present disclosure, the effect of interruptions in the
20 cardiac cycle and/or other irregularities in the data is further minimized and/or mitigated by monitoring the pressure differential calculations to detect these anomalies and automatically exclude the impacted cardiac cycles.

In one particular embodiment, pressure differential is calculated on two sequential cardiac cycles and the individual pressure differential values are averaged. The pressure
25 differential of a third cycle is then calculated. The average value of the pressure differentials is compared to the average pressure differential using three cycles. If the difference between the averages is below a predetermined threshold value, then the calculated value is considered to be stable and no further calculations are performed. For example, if a threshold value of 0.001 is used and adding an additional cardiac cycle changes the average pressure differential
30 value by less than 0.001, then the calculation is complete. However, if the difference between the averages is above the predetermined threshold value, then the pressure differential for a fourth cycle is calculated and a comparison to the threshold value is performed. This process is repeated iteratively until the difference between the averages of cardiac cycle N and cardiac cycle N+1 is below the predetermined threshold value. As the

pressure differential value is typically expressed to two decimal places of precision (such as 0.80), the threshold value for completing the analysis is typically selected to be small enough that adding a subsequent cardiac cycle will not change the pressure differential value. For example, in some instances the threshold value is selected to be between about 0.0001 and about 0.05.

In some instances, the level of confidence calculation has different thresholds depending on the degree of stenosis and/or an initial calculated pressure differential value. In that regard, pressure differential analysis of a stenosis is typically based around a cutoff value(s) for making decisions as to what type of therapy, if any, to administer. Accordingly, in some instances, it is desirable to be more accurate around these cutoff points. In other words, where the calculated pressure differential values are close to a cut-off, a higher degree of confidence is required. For example, if the cutoff for a treatment decision is at 0.80 and the initial calculated pressure differential measurement is between about 0.75 and about 0.85, then a higher degree of confidence is needed than if the initial calculated pressure differential measurement is 0.40, which is far from the 0.80 cutoff point. Accordingly, in some instances the threshold value is at least partially determined by the initial calculated pressure differential measurement. In some instances, the level of confidence or stability of the calculated pressure differential is visually indicated to user via a software interface.

Because pressure differential can be calculated based on a single cardiac cycle in accordance with the present disclosure, a real-time or live pressure differential calculation can made while the distal pressure measuring device is moved through the vessel. Accordingly, in some instances the system includes at least two modes: a single-cardiac-cycle mode that facilitates pressure differential calculations while moving the distal pressure measuring device through the vessel and a multi-cardiac-cycle mode that provides a more precise pressure differential calculation at a discrete location. In one embodiment of such a system, the interface 170 is configured to provide the live pressure differential value until the distal pressure measuring device is moved to the desired location and a measurement button is selected and/or some other actuation step is taken to trigger the multi-cardiac-cycle mode calculation.

Persons skilled in the art will also recognize that the apparatus, systems, and methods described above can be modified in various ways. Accordingly, persons of ordinary skill in the art will appreciate that the embodiments encompassed by the present disclosure are not limited to the particular exemplary embodiments described above. In that regard, although illustrative embodiments have been shown and described, a wide range of modification,

change, and substitution is contemplated in the foregoing disclosure. It is understood that such variations may be made to the foregoing without departing from the scope of the present disclosure. Accordingly, it is appropriate that the appended claims be construed broadly and in a manner consistent with the present disclosure.

CLAIMS

What is claimed is:

- 5 1. An interface for intravascular pressure sensing devices, comprising:
 - a distal input configured to receive a distal pressure signal from a distal pressure sensing device;
 - a distal output configured to output the distal pressure signal to a hemodynamic system in a format useable by the hemodynamic system;
 - 10 a proximal input configured to receive a proximal pressure signal from a proximal pressure sensing device;
 - a proximal output configured to output the proximal pressure signal to the hemodynamic system in a format useable by the hemodynamic system;
 - a processor coupled to the distal input, distal output, proximal input, and proximal output, the processor configured to calculate a pressure differential between the distal pressure and the proximal pressure based on the received distal pressure signal and the received proximal pressure signal; and
 - 15 a display in communication with the processor and configured to display the pressure differential calculated by the processor.
- 20 2. The interface of claim 1, wherein the distal input, distal output, proximal input, proximal output, processor, and display are secured to a housing.
3. The interface of claim 2, wherein the housing has a width between 5 cm and 25 cm, a height between 5 cm and 25 cm, and a depth between 1 cm and 10 cm.
- 25 4. The interface of claim 3, wherein the distal pressure sensing device is a pressure-sensing guidewire.
- 30 5. The interface of claim 4, wherein the proximal pressure sensing device is a pressure-sensing catheter configured for use with the hemodynamic system.
6. The interface of claim 5, wherein the pressure-sensing catheter communicates with the hemodynamic system via at least four leads.

7. The interface of claim 6, wherein a first pair of leads of the at least four leads are associated with sending an excitation signal to the pressure-sensing catheter.

5 8. The interface of claim 7, further comprising a first amplifier electrically coupled with the first pair of leads, the first amplifier configured to sample the excitation signal.

9. The interface of claim 8, wherein the first amplifier sends the sampled excitation signal to the microprocessor.

10

10. The interface of claim 9, wherein a second pair of leads of the at least four leads are associated with sending the proximal pressure signal from the proximal pressure sensing device to the hemodynamic system.

15 11. The interface of claim 10, further comprising a second amplifier electrically coupled with the second pair of leads, the second amplifier configured to sample the proximal pressure signal.

20 12. The interface of claim 11, wherein the second amplifier sends the sampled proximal pressure signal to the microprocessor.

25 13. The interface of claim 1, further comprising a power extraction device coupled to a pair of leads that receive an excitation signal from the hemodynamic system, the power extraction device positioned within the housing and configured to extract power from the excitation signal for use in operating at least the processor and the display.

14. The interface of claim 13, wherein the pair of leads are associated with a proximal pressure measurement input of the hemodynamic system.

30 15. The interface of claim 13, wherein the pair of leads are associated with a distal pressure measurement input of the hemodynamic system.

16. The interface of claim 1, wherein the processor is further configured to identify a diagnostic window for calculating the pressure differential, the diagnostic window consisting of a portion of a heartbeat cycle of a patient.

5 17. A system for evaluating a vascular stenosis, the system comprising:
a distal pressure sensing device sized and shaped for insertion into human vasculature;
a proximal pressure sensing device sized and shaped for insertion into human
vasculature; and

an interface comprising:

10 a distal input configured to receive a distal pressure signal from the distal
pressure sensing device;

a proximal input configured to receive a proximal pressure signal from the
proximal pressure sensing device;

15 a proximal output configured to output the proximal pressure signal to a
processing system in a format useable by the processing system;

a processor coupled to the distal input, proximal input, and proximal output,
the processor configured to calculate a pressure differential between the distal
pressure and the proximal pressure based on the received distal pressure signal and
the received proximal pressure signal; and

20 a display in communication with the processor and configured to display the
pressure differential calculated by the processor.

18. The system of claim 17, wherein the processing system is a hemodynamic system.

25 19. The system of claim 17, wherein the display is a touchscreen.

20. The system of claim 17, wherein the display is further configured to display the
proximal pressure and the distal pressure.

30 21. The system of claim 17, wherein the interface further comprises a distal output
configured to output the distal pressure signal to a processing system in a format useable by
the processing system.

22. The system of claim 21, wherein the distal input, distal output, proximal input, proximal output, processor, and display are secured to a housing.

23. The system of claim 22, wherein the housing has a width between 5 cm and 25 cm, a
5 height between 5 cm and 25 cm, and a depth between 1 cm and 10 cm.

24. The system of claim 23, wherein the distal pressure sensing device is a pressure-sensing guidewire.

10 25. The system of claim 24, wherein the proximal pressure sensing device is a pressure-sensing catheter configured for use with the processing system.

26. The system of claim 25, wherein the pressure-sensing catheter communicates with the processing system via at least four leads.

15

27. The system of claim 26, wherein a first pair of leads of the at least four leads are associated with sending an excitation signal to the pressure-sensing catheter.

28. The system of claim 27, further comprising a first amplifier electrically coupled with
20 the first pair of leads, the first amplifier configured to sample the excitation signal.

29. The system of claim 28, wherein the first amplifier sends the sampled excitation signal to the microprocessor.

25 30. The system of claim 29, wherein a second pair of leads of the at least four leads are associated with sending the proximal pressure signal from the proximal pressure sensing device to the processing system.

31. The system of claim 30, further comprising a second amplifier electrically coupled
30 with the second pair of leads, the second amplifier configured to sample the proximal pressure signal.

32. The system of claim 31, wherein the second amplifier sends the sampled proximal pressure signal to the microprocessor.

33. The system of claim 17, further comprising a power extraction device coupled to a pair of leads that receive an excitation signal from the processing system, the power extraction device positioned within the housing and configured to extract power from the excitation signal for use in operating at least the processor and the display.
34. The system of claim 33, wherein the pair of leads are associated with a proximal pressure measurement input of the processing system.
35. The system of claim 33, wherein the pair of leads are associated with a distal pressure measurement input of the processing system.
36. The system of claim 17, wherein the processor is further configured to identify a diagnostic window for calculating the pressure differential, the diagnostic window consisting of a portion of a heartbeat cycle of a patient.

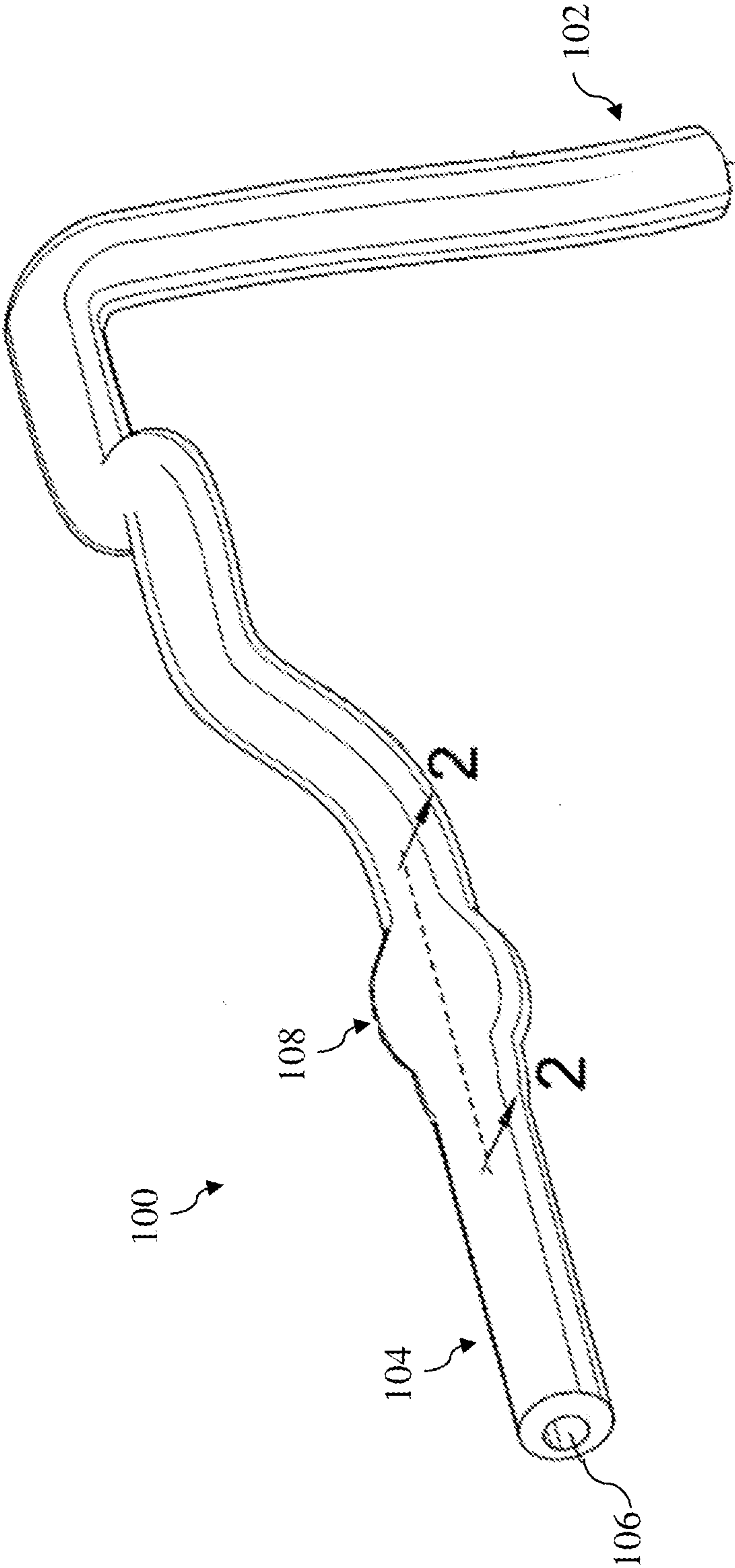


Fig. 1

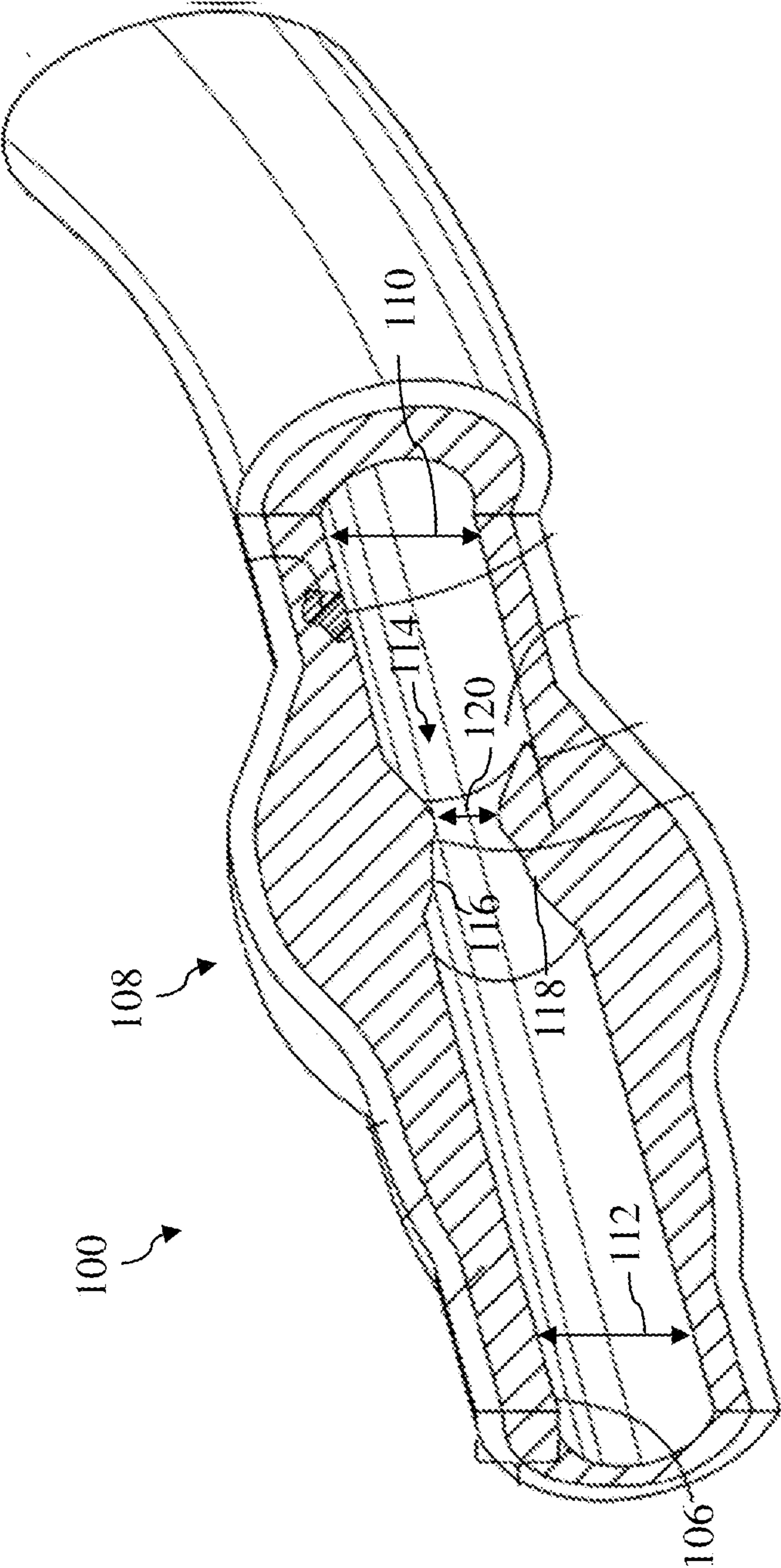


Fig. 2

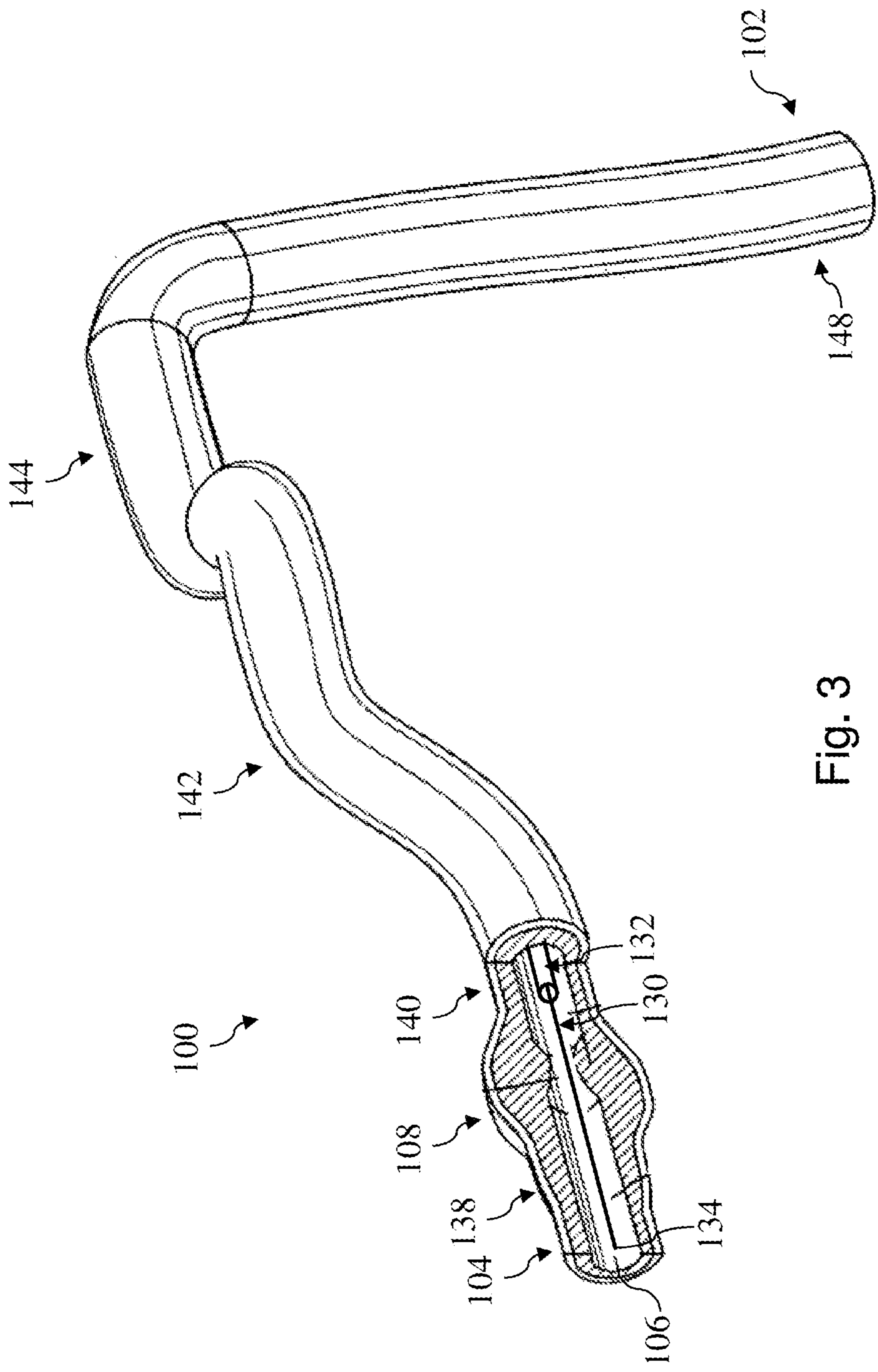


Fig. 3

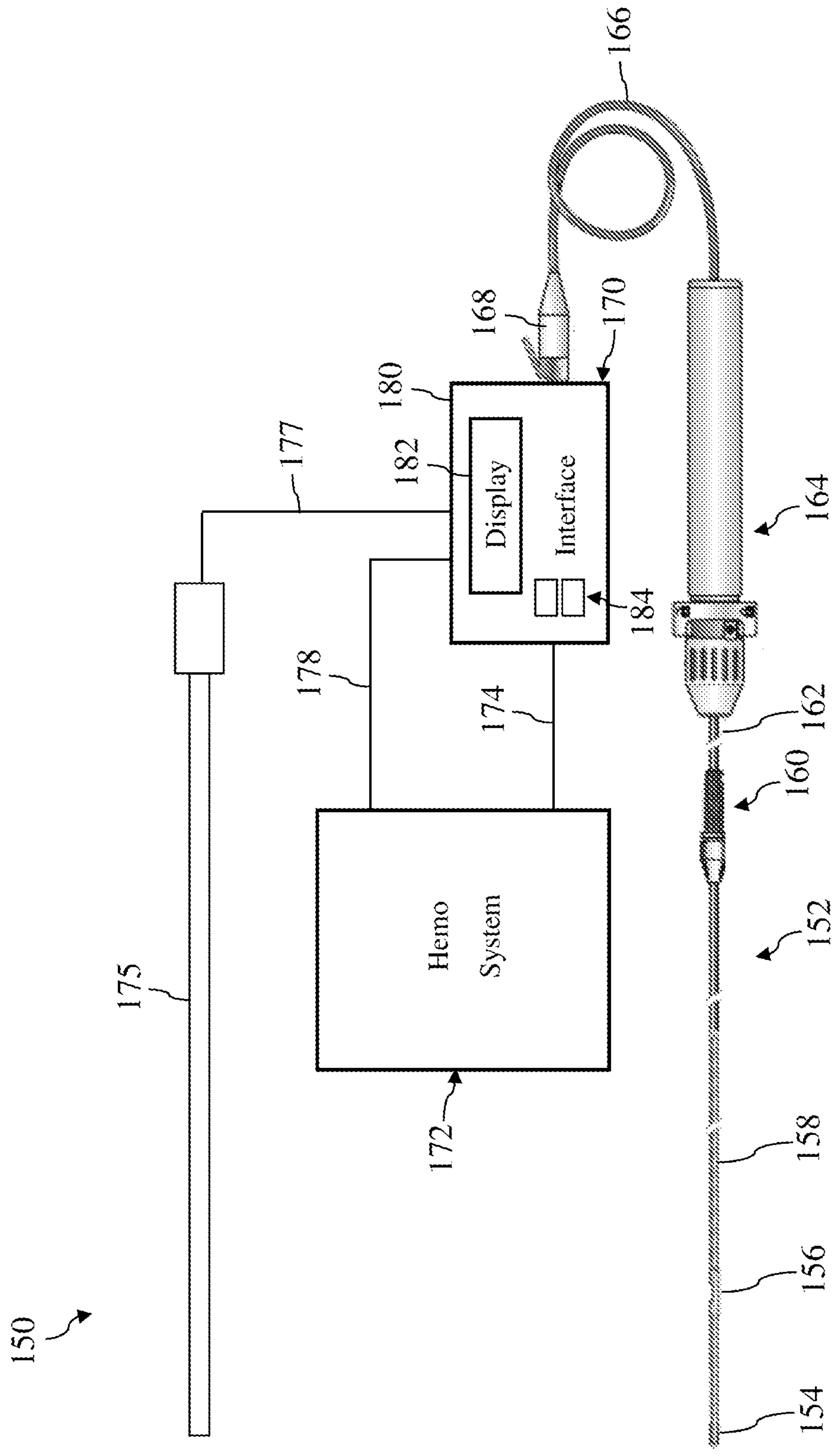


Fig. 4

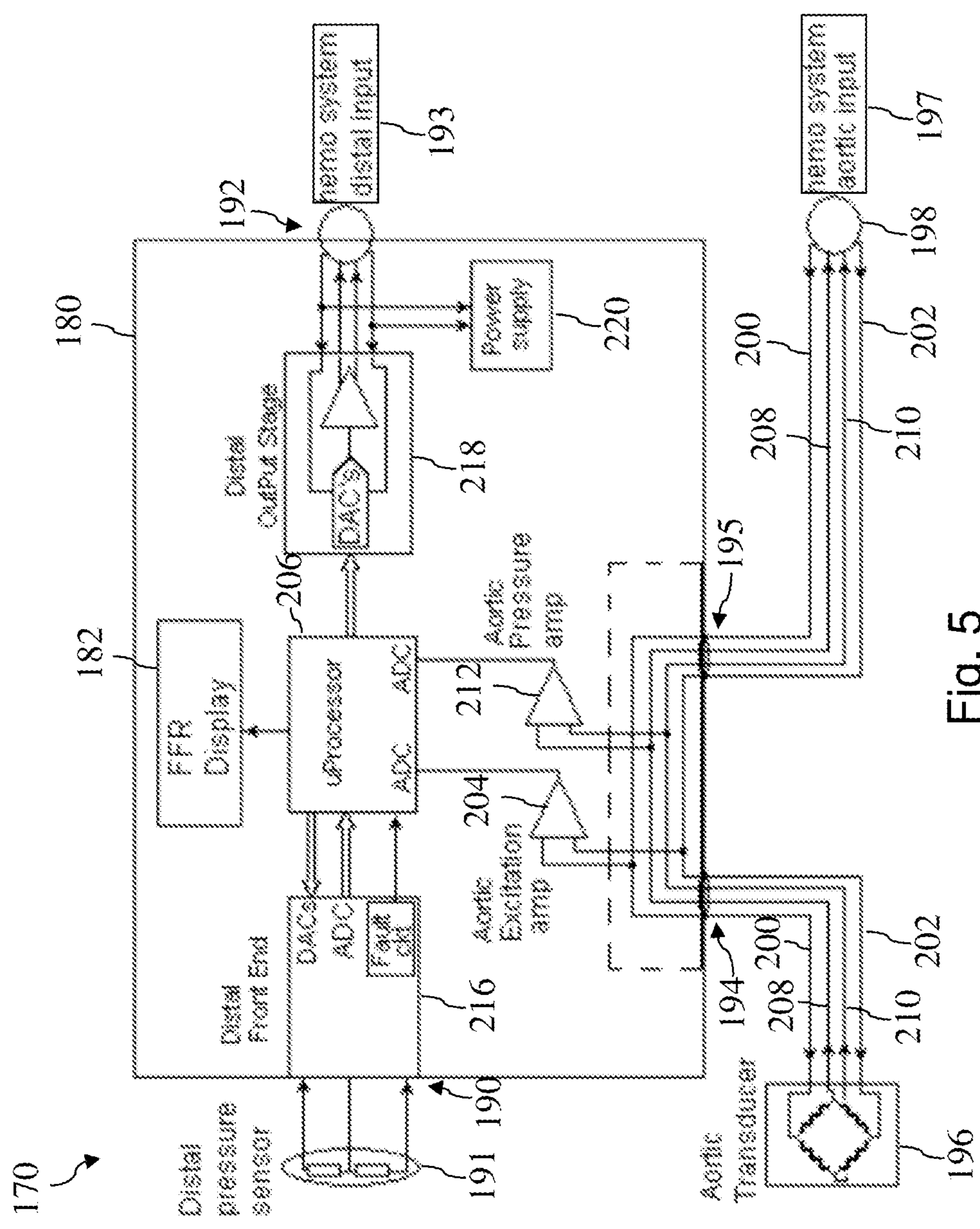


Fig. 5

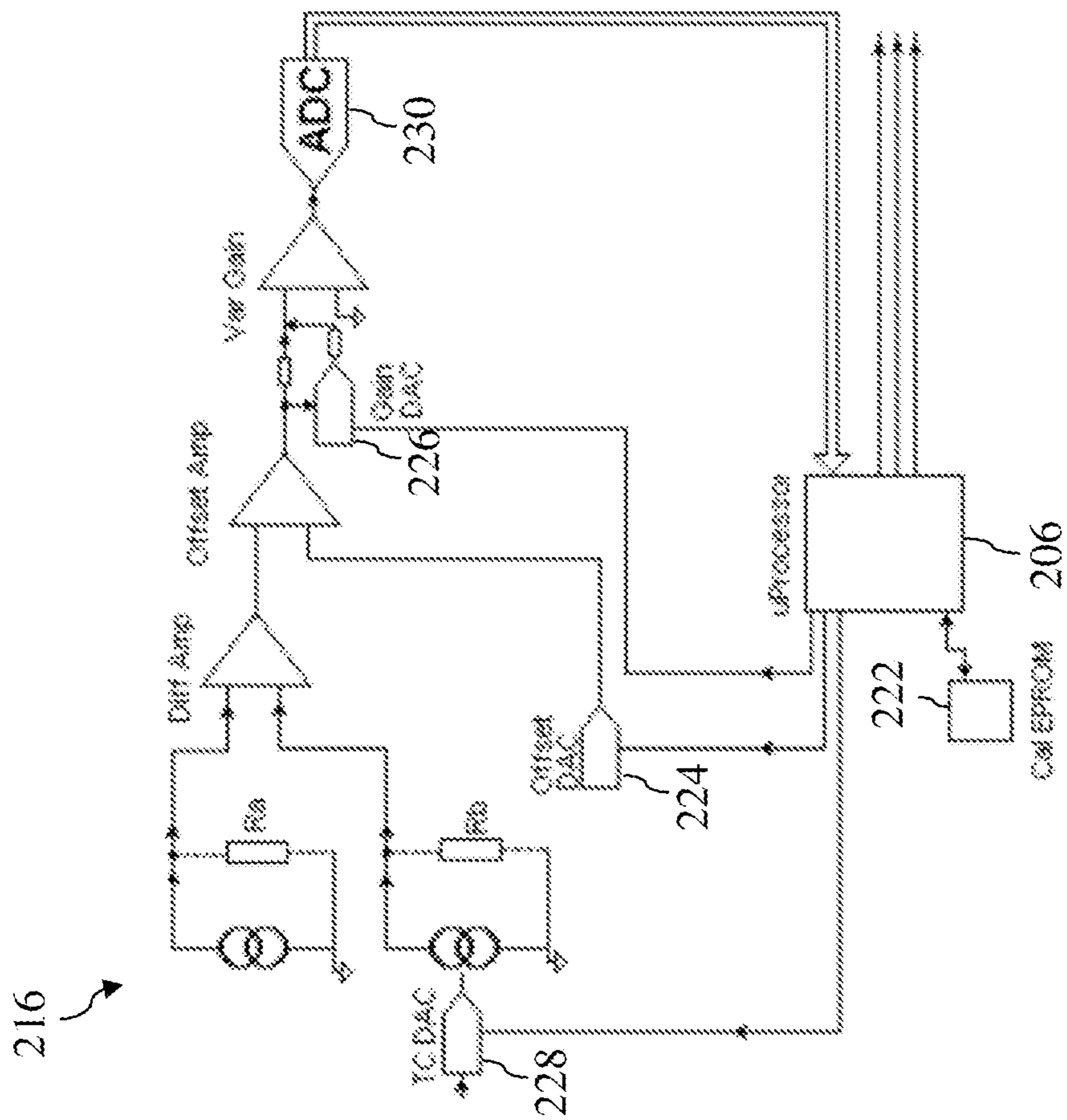


Fig. 6

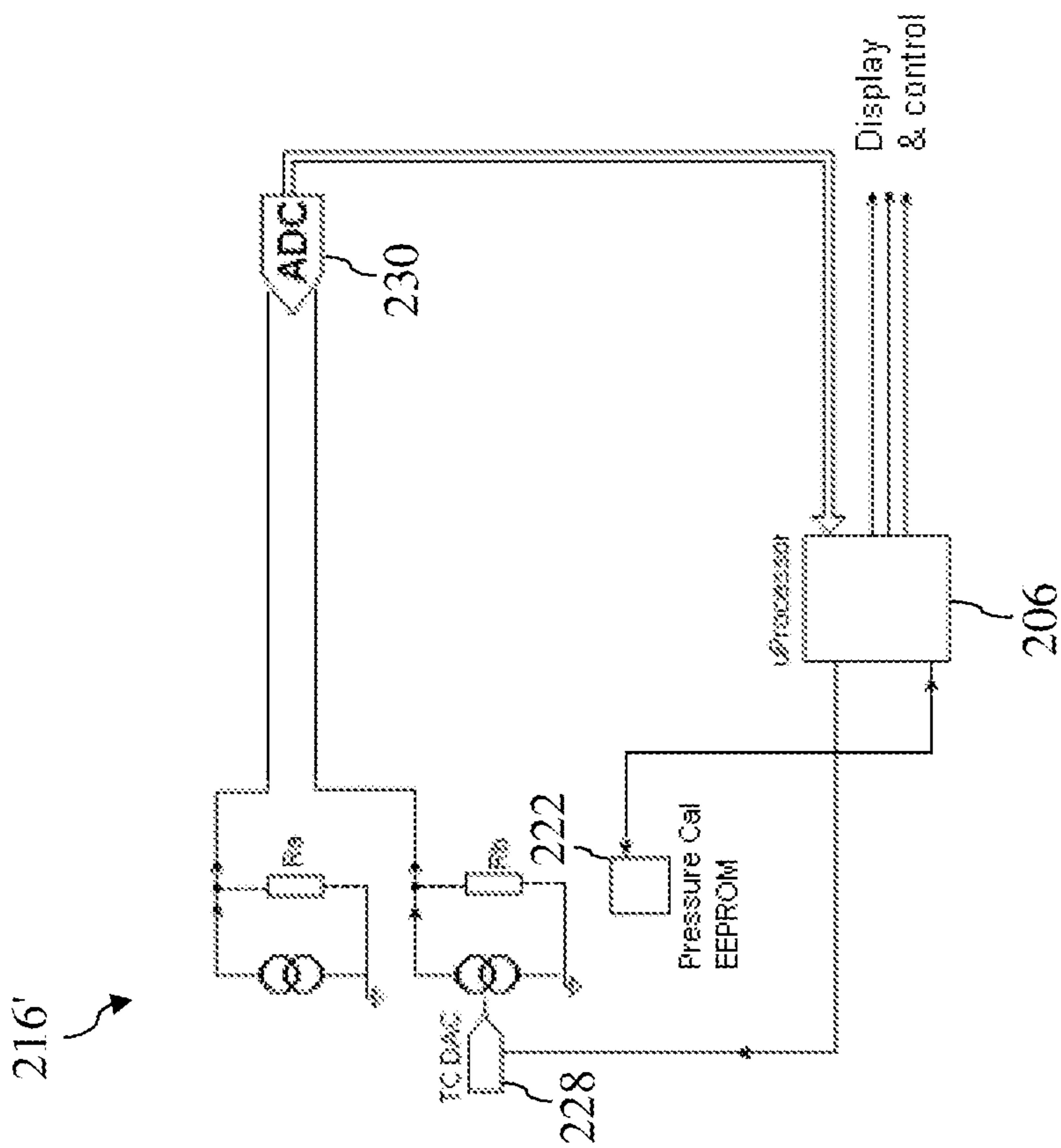


Fig. 7

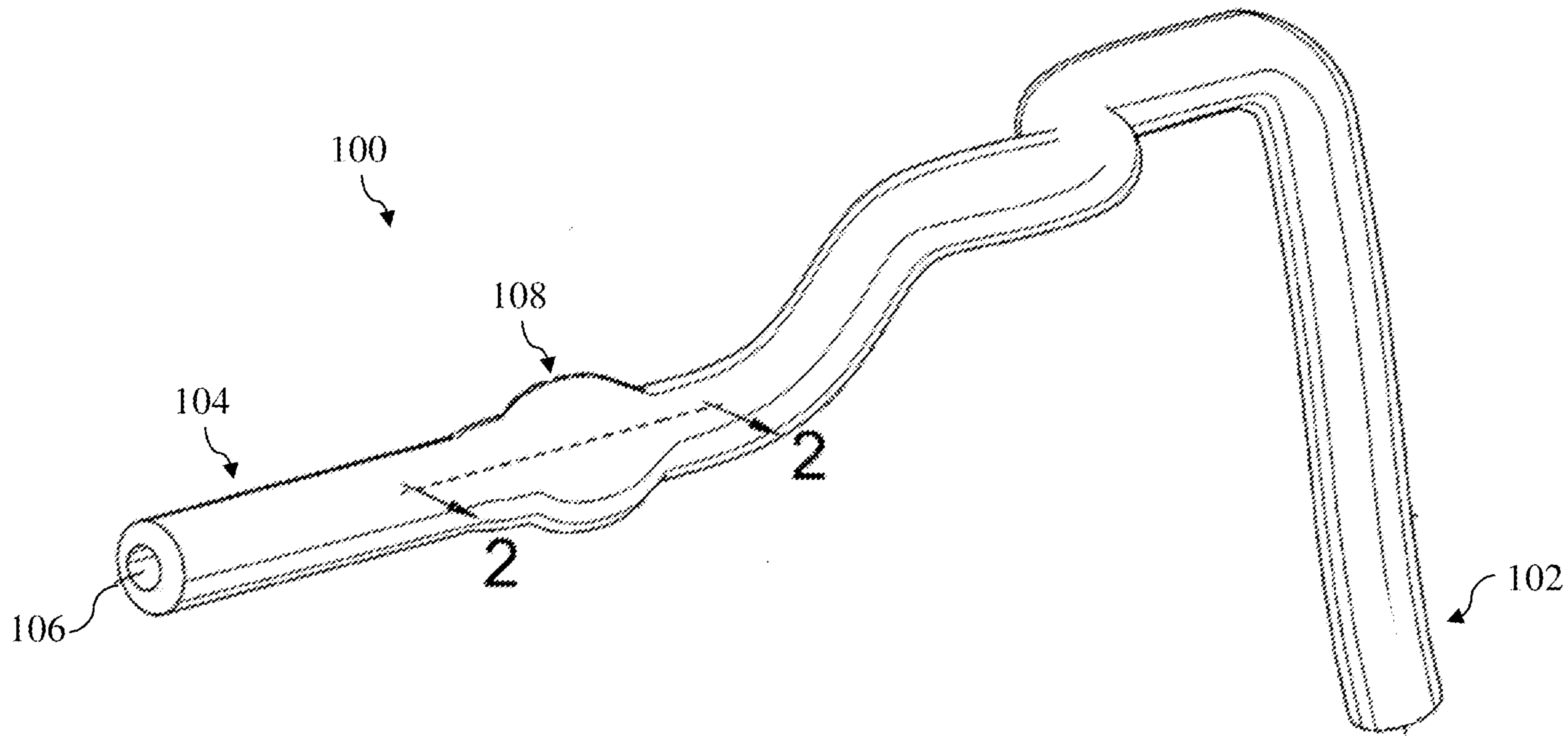


Fig. 1